



## ***Cordyceps locustiphila* (Hypocreales: Cordycipitaceae) infecting the grasshopper pest *Tropidacris collaris* (Orthoptera: Acridoidea: Romaleidae)**

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With 3 figures

**Abstract:** *Cordyceps locustiphila* is described, illustrated, and compared with a previous finding. This species was collected infecting *Tropidacris collaris* in the Yabotí biosphere reserve, Misiones province, Argentina. Until now, this entomopathogenic fungus has not been reported affecting *T. collaris*, and it is the first report for Argentina. From the cordyceps-like stromata of *C. locustiphila* it was possible to isolate for the first time the acremonium-like anamorph of this fungus. The identity of both, anamorph and teleomorph, was determined by morphological and molecular taxonomic studies, which challenges the recent new combination into *Beauveria*.

**Key words:** entomogenous fungi, insect pathogens, tropical biodiversity.

### **Introduction**

Fungi with cordyceps-like teleomorph are most diverse in the family *Cordycipitaceae* in terms of both number of known species and host range (Kobayasi 1941, Sung et al. 2007a). There are estimated to be more than 400 species names under *Cordyceps* (Mains 1958, Stensrud et al. 2005) although this is expected to be an underestimation of the

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extant global diversity (Hawksworth & Rossman 1997). A phylogenetic classification based on molecular data separated the species across three families, *Cordycipitaceae*, *Clavicipitaceae* and *Ophiocordycipitaceae* (Sung et al. 2007b, Sanjuan et al. 2014). Cordyceps-like fungi have the highest species diversity in subtropical and tropical regions (Samson et al. 1988) such as tropical areas of Brazil (Andrade 1980, Evans & Samson 1982, Evans et al. 2011), Colombia (Kobayasi 1981, Sanjuan et al. 2001, 2014, 2015) Ecuador (Kobayasi 1981, Evans & Samson 1982), and Bolivia (Mains 1959). In Argentina, these fungi had been found infecting different insects (Spegazzini 1919, Marchionatto 1945, Mains 1954, 1959, Yasem de Romero 1984, López Lastra 1989, Mueller & Rajchenberg 1991). *Cordyceps locustiphila* was described by Hennings (1904) as a species with gregarious or solitary, claviform, yellow stromata. The original illustration shows the fungus emerging from the abdomen and the coxa of an adult locust (Orthoptera: Acrididae). For molecular phylogenetic and nomenclatural reasons, this species was transferred to *Beauveria* (Kepler et al. 2017).

The romaleid *Tropidacris collaris* (Stoll) (Orthoptera: Acridoidea: Romaleidae), one of the largest grasshoppers known ( $\sigma = 73\text{--}101$  mm,  $\text{♀} = 92\text{--}126$  mm), has become in recent years a recurrent and extended pest in areas of some of the northern provinces of Argentina (Catamarca, Chaco, Córdoba, La Rioja, Santiago del Estero), particularly in olive and jojoba plantations (Cigliano et al. 2014).

Until now, except for an undescribed *Gregarina* sp. (Bardi et al. 2009), pathogens have not been reported affecting *T. collaris*. Here we first report for Argentina by morphological and molecular studies an isolate of *C. locustiphila* from *T. collaris*. Moreover, from stromata of *C. locustiphila* we were able to isolate for the first time the asexual phase of this fungus, whose characters, were studied with morphological observations and molecular techniques.

## Materials and methods

**FIELD COLLECTION:** The specimen of *T. collaris* that was infected by *C. locustiphila* was found in the Yabotí biosphere reserve (26°37'S, 53°40'W), a natural protected area that covers parts of the departments of Guaraní and San Pedro in Misiones province, Argentina. The area presents an extension of 235,959 ha and belongs to the Selva Paranaense (Parana forest) eco-region (Cabrera & Willink 1973, Morrone 2014). The climate is subtropical without dry season (Cabrera & Willink 1973, Di Bitetti et al. 2003). Average temperatures range from 24°C in summer to 14°C in winter. The rainfall is 1800 mm annually, concentrated in the summer season. This reserve has an exuberant jungle, made up of lianas, epiphytic plants, ferns, and tree species such as black laurel [*Nectandra saligna* Nees & Mart.], white guatambú [*Balfourodendron riedelianum* (Engl.) Engl.], yerba mate (*Ilex paraguariensis* A. St.-Hil.), and guayubira (*Patagonula americana* L.), among many others. The grasshopper *T. collaris* was collected in January 2015 in a region within the reserve that was not altered by human activity, where there are recorded temperatures of 24°C and an ambient humidity of 90%. The grasshopper *T. collaris* infected with *C. locustiphila* was placed in a plastic box with silica gel and deposited at the Herbarium of the Spegazzini Institute of Botany of La Plata National University, with access code LPS 49245.

**MORPHOLOGICAL OBSERVATIONS:** In the laboratory, fungal stromata were taken from the infected insect. They were surface-sterilized by dipping them successively in 70% ethanol (10–15 s), 0.5% sodium hypochlorite solution (1 min), and sterile distilled water (1 min, two consecutive baths) according to Lacey & Solter (2012). Stromata were allowed to dry at room temperature under a laminar flow



Fig. 1. Stromata of *Cordyceps locustiphila* growing on *Tropidacris collaris* grasshopper. Bars: A = 6 mm; B = 4 mm.

chamber. Then each stroma was cut longitudinally into two equal halves with a sterile scalpel, and deposited on a Petri dish containing potato dextrose agar (PDA) plus antibiotics (0.1% stock antibiotics consisting of 0.02 g each of tetracycline, streptomycin and penicillin) as culture medium. The plates were then incubated at 25°C for 7 days, after which fungal isolates were transferred to fresh PDA. A culture was deposited in the living strain collection of LPS under the number LPSc #1218. Some stromata were fixed with a Bouin's solution and prepared (Becnel 2012) to perform histological sections for observation of stipes, perithecia, ascospores, and asci under stereoscopic and optical microscopes with phase contrast to corroborate the taxonomic identification using available keys and databases (Kobayasi 1941, 1981, Luangsa-ard et al. 2008).

**DNA EXTRACTION, PCR, AND SEQUENCING:** To confirm the morphological identification, a portion of the stroma and cultivated mycelium were placed separately in CTAB buffer to perform the extraction of total DNA following the procedure described by Stenglein & Balatti (2006), then amplifications were performed by the Polymerase Chain Reaction (PCR). The regions corresponding to the internal transcriber spacer region (ITS) and to the elongation factor 1 $\alpha$  gene (TEF) were amplified according to Sanjuan et al. (2014), using primers ITS4 / ITS 5 (White et al. 1990) and 983F / 2218R (Rehner & Buckley 2005), respectively. Finally, the samples obtained were sequenced and deposited at GenBank (accession numbers: MF185185; MF185186; MF185187; MF185188). The sequences were submitted to BLAST searches at GenBank.

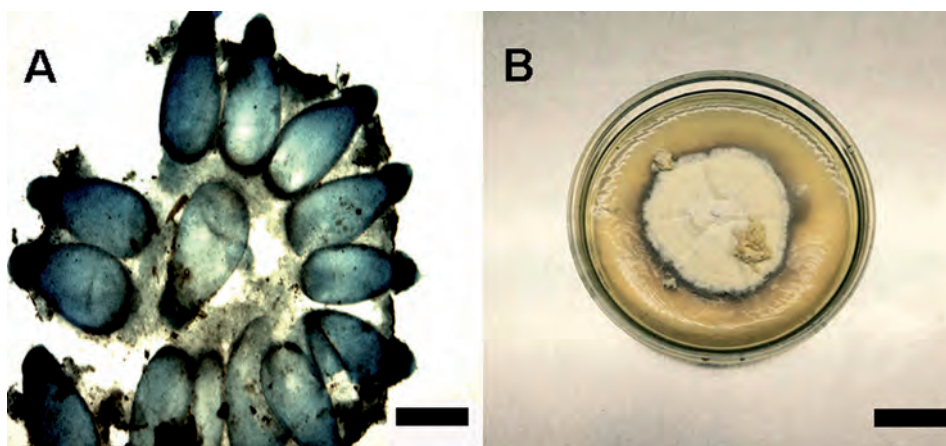


Fig. 2. A: Perithecia (in cotton blue) of *Cordyceps locustiphila*. B: Colony on PDA after 15 days at 25°C. Bars: A = 150  $\mu$ m; B = 15 mm.

## Results

### Taxonomic description

*Cordyceps locustiphila* Henn., Fungi amazonici II, a cl. Ernesto Ule collecti. Hedwigia 43: 246 (1904).

Teleomorph characteristics (LPS#49245): Stromata gregarious, claviform, simple, bright yellow (Fig. 1A–B), 5–10 mm long. Fertile head clavate, slightly echinulate by protruding perithecial ostioles, bright yellow, 3–5  $\times$  2–4 mm long. Stipe fleshy, terete, sometimes caespitose, grayish yellow, 1–4  $\times$  1–2 mm. Perithecia semi-immersed, with perpendicular orientation, ovoid, 358–488  $\times$  138–232  $\mu$ m ( $n = 20$ ), wall less than 50  $\mu$ m wide (Fig. 2A). We did not observe the presence of asci and ascospores, because stromata of *C. locustiphila* did not mature, even after placing them in humid chamber. ITS (accession number: MF185185) and TEF (accession number: MF185186) sequences obtained from the culture mycelium showed 100% similarity with GenBank accessions JQ958609 and JQ958619 of *C. locustiphila*, respectively.

### Anamorph characteristic

Colonies obtained from stroma tissue slow-growing, after 15 days on PDA at 25°C 30 mm diam., ocher yellow, white near margin; reverse pale yellow (Fig. 2B). Hyphae 1.5–2.5 ( $\pm 0.08$ ) ( $n = 50$ )  $\mu$ m wide. Phialides acromonium-like, simple, awl-shaped, erect, sessile phialides from the substrate 40.6–56.8 ( $\pm 5.38$ )  $\times$  1.1–2.1 ( $\pm 0.34$ )  $\mu$ m ( $n = 50$ ) wide at the base and 0.71–1.25 ( $\pm 0.18$ )  $\mu$ m wide at the tip ( $n = 50$ ) (Fig. 3). Conidia hyaline, smooth, cylindrical, slightly clavate or allantoid, 1.2–3.1 ( $\pm 0.63$ )  $\times$  4.7–9.8 ( $\pm 1.79$ )  $\mu$ m ( $n = 100$ ), always arranged in slime drops or solitary (Fig. 3). Sequences of the anamorph in culture differed only at bp position 391 (C/G) for the

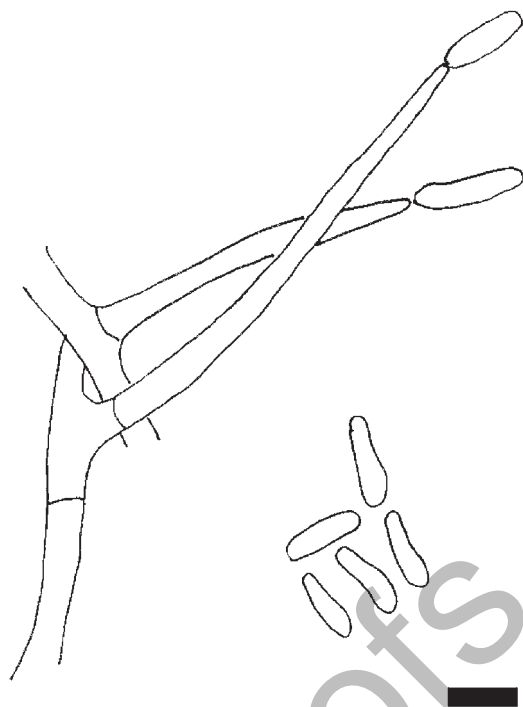


Fig. 3. Hypha, phialides and conidia of *Cordyceps locustiphila*. Bars = 5  $\mu$ m.

ITS region, and at bp 213 (A/C) and 221 (T/A) for the TEF region to those derived from the ascostroma.

SPECIMEN EXAMINED: ARGENTINA. MISIONES, Departamento, Yabotí biosphere reserve: 26°37'S, 54°10'W, 200 to 648 m.s.n.m., on *Tropidacris collaris* (Orthoptera: Acridoidea: Romaleidae), (LPSC WDCM1001- La Plata Spegazzini Collection; and [www.museo.fcnym.unlp.edu.ar/micologia\\_colecciones](http://www.museo.fcnym.unlp.edu.ar/micologia_colecciones)), S. Pelizza, LPS#49245, deposited in the herbarium of the Spegazzini Institute in November 2015, culture LPSc 1218.

## Discussion

Our detection constitutes the first record of *C. locustiphila* infecting *T. collaris*. Until now *C. locustiphila* was known to occur only in the grasshopper *Colpolopha* sp., also among the Romaleidae, the most diversified family of endemic neotropical grasshoppers (Cigliano et al. 2014). This record of *C. locustiphila* is the first for Argentina and the southernmost known one, since the previous records of this entomopathogenic fungus were carried out in different regions of the Amazon jungle in Brazil, Colombia, Ecuador, and Perú (Sanjuan et al. 2014).



Our specimen appears related with the one described by Sanjuan et al. (2014) parasitizing *Colpolopha laetipenis* due to the gregarious, claviform, simple, and bright yellow stromata, but differs from it in size. In Sanjuan et al. (2014) the stromata were 12–20 mm long while in our specimen stromata were 5–10 mm long. Differences between these two specimens of *C. locustiphila* are also found in perithecia. The perithecia of *C. locustiphila* found in *Colpolopha* measured 550–600 x 250–320 µm, whereas those found on *T. collaris* measured 358–488 x 138–232 µm. In summary, we observed that the different structures measured (stromata and perithecia) were slightly smaller in *C. locustiphila* found on *T. collaris* than that found in specimens of the genus *Colpolopha*., which may be based on the immaturity of our specimen.

Until now, it was speculated that *C. locustiphila* had a *beauveria* anamorph (Sanjuan et al. 2014), but this speculation could not be corroborated. The molecular analysis on the acremonium-like anamorph obtained in culture strongly confirmed that it belongs to *C. locustiphila*. The recent new combination *B. locustiphila* was made merely to comply with the 1F = 1N rule without providing new data for this species (Kepler et al. 2017). Our finding of an acremonium-like anamorph indicates that this combination might be premature and that more study is necessary in the *Beauveria* clade before sound nomenclatural conclusions can be proposed.

Even though further research is necessary in order to properly describe the anamorph we think it is important to communicate the detection of this new isolate. When outbreaks of *T. collaris* occur, control measures still rely exclusively on chemical insecticides and the eventual development of a biological control alternative would be a significant environmental-friendly step forward.

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